Verdibasert helsetjeneste: fra teori til virkelighet

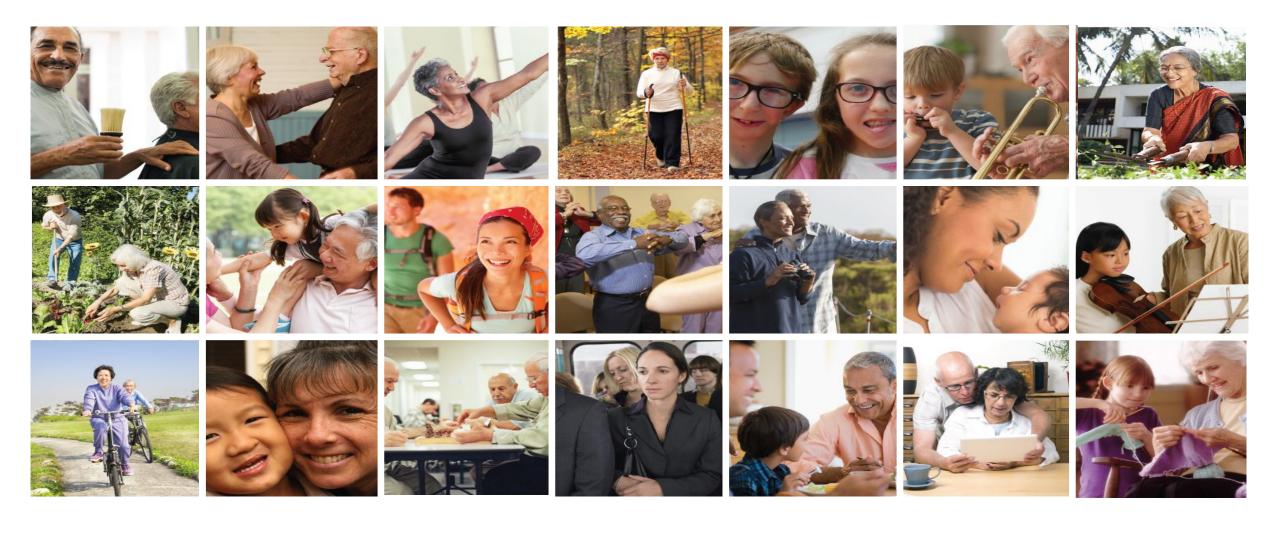
Del I. Moving towards an outcome- and value-based environment

Dr. Christina R Åkerman

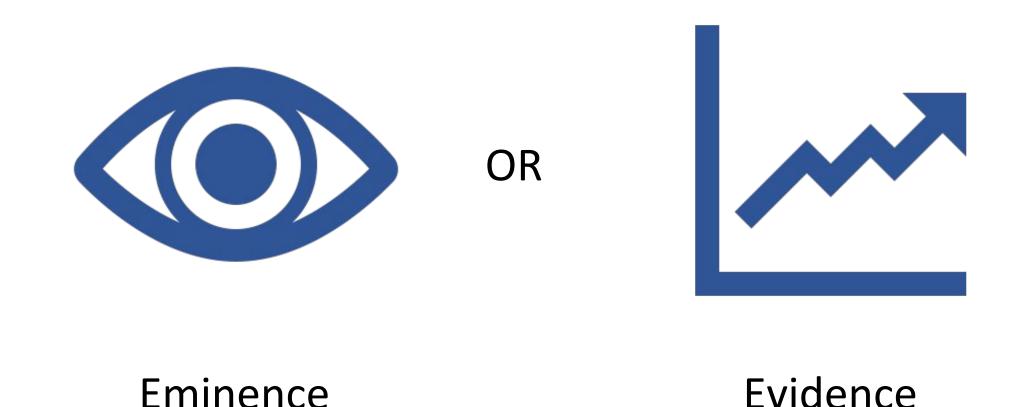
Nov 5, 2019

HelseCampus Stavanger & Norwegian Smart Care Cluster

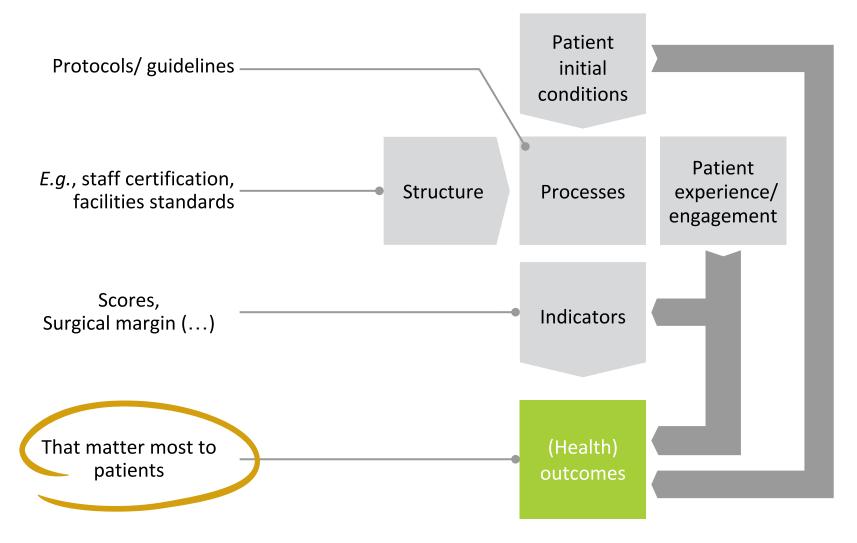
Putting the focus on the patient



Generally speaking, what do we primarily base our judgements about health care services on?

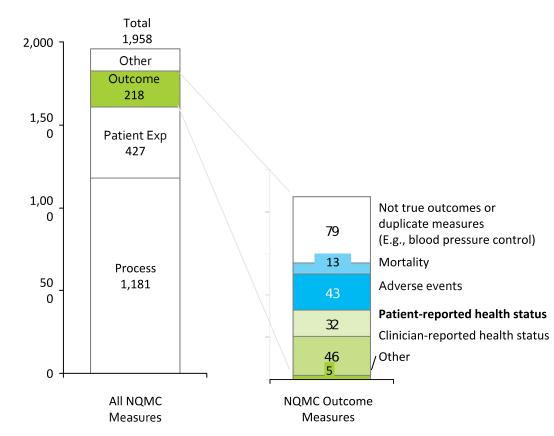


Outcomes are the "real-world" results that matter most to patients



Moving to value-based health care demands a better way of measuring outcomes

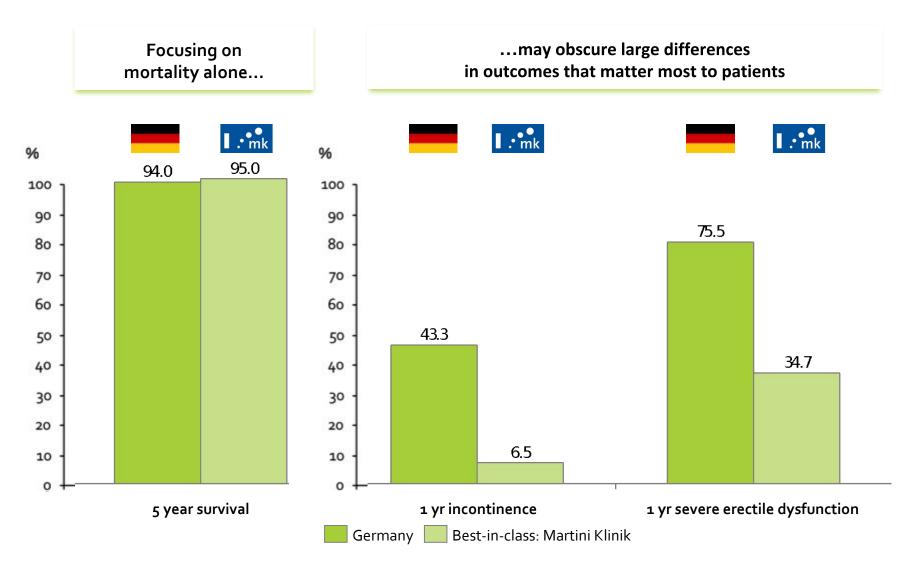
Outcomes, especially patient-reported outcomes are significantly under-represented in available measures¹



NQMC: National Quality Measures Clearinghouse

Measuring meaningful outcomes matter

Comparing outcomes of prostate cancer care



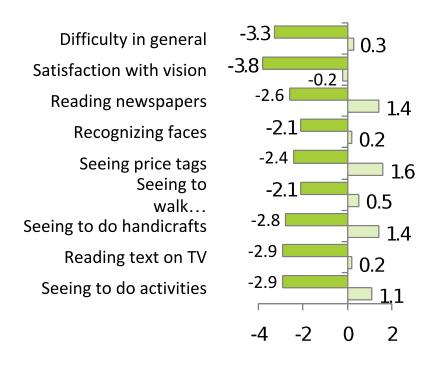
Why the patient's perspective matters

Study

- Analysis of clinical outcomes and patientreported outcomes in Swedish Cataract Registry
 - 9,707 patients
- Of those, nearly all those surveyed showed clinical outcome improvement (VA)
 - 97.8%
- Interestingly, of those 97.8%, 7.4% reported being worse off visually than before

How is this possible?

Findings





"Every hospital should follow every patient it treats long enough to determine whether or not the treatment has been successful, and then to inquire, 'if not, why not?' with a view to preventing similar failures in the future"

- Ernest Codman (1914, Boston)

Redefining Health Care

Michael E Porter & Elizabeth Olmsted Teisberg 2006

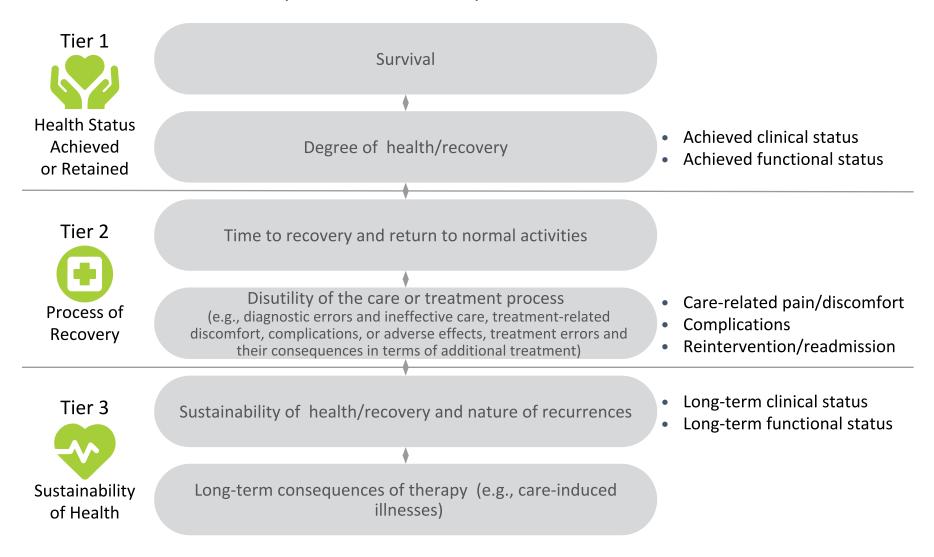
Patient health outcomes achieved

Value =

Cost of delivering those outcomes

The Outcome Measures Hierarchy

Two Dimensions- Clinician Reported and Patient Reported Outcomes



Source: NEJM Dec 2010

International Consortium for Health Outcomes Measurement ICHOM

Principles of Standard Set Development

- Outcomes are **defined around medical conditions**, not specialties or the procedures
- Standard Sets are a "minimum set" focused on outcomes that matter most to patients
- Patients are directly involved in defining every Standard Set

Patient-reported outcomes are part of every

- Standard Set and include functional status, symptom burden and health-related quality of life
- A "minimum set" of initial conditions/risk factors is included to facilitate meaningful comparison
- Time points and sources of data collection are clearly defined to ensure comparability of results

"GLOBAL LANGUAGE FOR HEALTH OUTCOMES"

ICHOM Standard Sets cover >50% of global disease burden

- 1. Localized Prostate Cancer
- 2. Low Back Pain
- 3. Coronary Artery Disease
- 4. Cataracts
- 5. Parkinson's Disease
- 6. Cleft Lip and Palate
- 7. Stroke
- 8. Hip and Knee Osteoarthritis
- 9. Macular Degeneration
- 10. Lung Cancer
- 11. Depression and Anxiety
- 12. Advanced Prostate Cancer
- 13. Heart Failure
- 14. Pregnancy and Childbirth

- 15. Colorectal Cancer
- 16. Older Person
- 17. Overactive Bladder
- 18. Craniofacial Microsomia
- 19. Inflammatory Bowel Disease
- 20. Chronic Kidney Disease
- 21. Hypertension
- 22. Inflammatory arthritis
- 23. Congenital upper limb anomalies
- 24. Pediatric facial palsy
- 25. Diabetes (I+II)
- 26. Atrial Fibrillation
- 27. Breast Cancer
- 28. Dementia

The Breast Cancer Standard Set Flyer represents a high-level overview of the outcomes, scope and treatments



Scope

All patients (men and women) with newly pathologically diagnosed invasive breast cancer (stage I-IV) and DCIS

Exclude:

- Rare tumor (e.g. Phyllodes tumor)
- Lobular carcinoma in situ (LCIS)
- Patients with recurrent disease at baseline

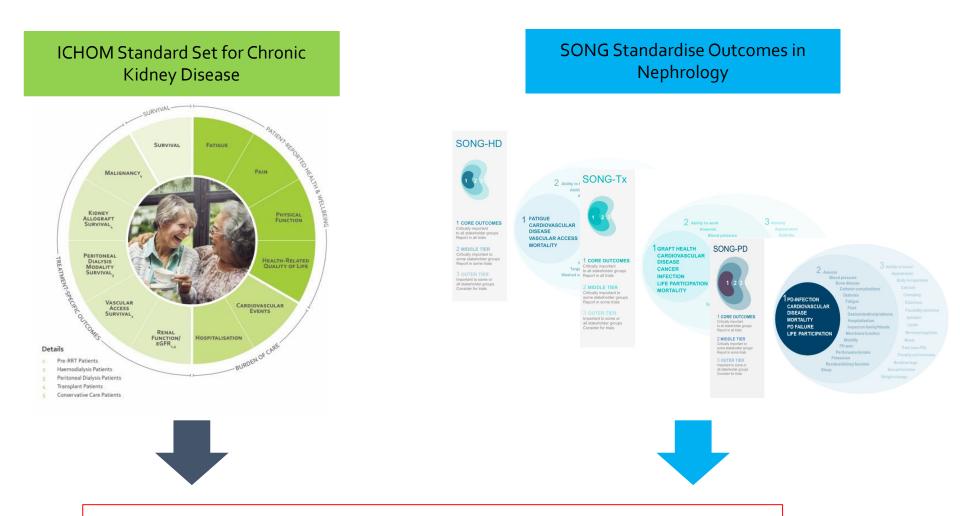
Treatment Approaches covered

- Surgery
- Radiotherapy
- Chemotherapy
- Targeted therapy
- Hormonal therapy

Exclude:

Investigational agents or techniques

Overlap in core outcome sets in clinical practise (ICHOM) and clinical trials (SONG)



Fatigue, cardiovascular disease, mortality, vascular access, PD-failure, graft survival

Driving the value of hospitals and service delivery: an OECD perspective IHF 2018 Volume 54 Number 3

A core objective of a healthcare organisation is to **maximise the quality of care for every patient**, but data on key quality dimensions of safety, effectiveness and people-centredness are **not systematically captured** from the patient's perspective.

This means that governing bodies are basing decisions that determine success in a competitive marketplace on **incomplete information**.

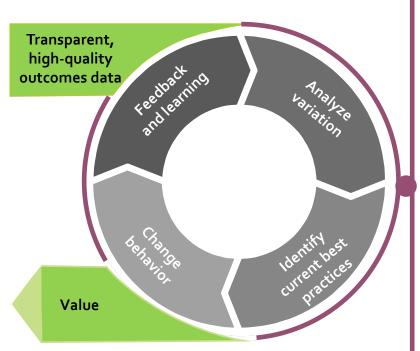
Addressing this requires routine measurement of outcomes and experiences from the patients themselves.

The **OECD's Paris** initiative is helping to build the capacity of countries and organisations to capture the voice of the patient through validated, comparable indicators, but successful implementation means engaging front-line staff and patients, and integrating these metrics into existing information infrastructure.

EU Health Summit Brussels Nov 29, 2018 "A shared vison for the future of health in Europe"



Outcome measurement empowers stakeholders to generate value



Key stakeholders



 Patients will choose their provider based on expected outcomes and their share of the cost



 Clinicians will improve quality of care by comparing performance and learning from each other



 Hospitals will differentiate into areas where they deliver superior outcomes at competitive prices



 Payers will negotiate contracts based on results, not volume, and encourage innovation to achieve those results



 Life science will market their products on value, showing improved outcomes relative to costs